Commonwealth of Virginia Department of Health Professions 6603 Board Street Road, 5th Floor Richmond, Virginia 23230

Date	
Hours	
Rev: 4/2005	

WAREHOUSE INSPECTION REPORT

Name				License No	Exp Date	<u></u>	
Street			City		_Zip		
Telephone No)		Fax No	Hours of Operation	n		
Responsible P	arty						
CSRC Licens	e No		Exp Date	DEA No	Exp	Date	
TYPE OF FA	CILIT	Y					
WAREHOUS	E 🗆	Presci	ription Drugs 🔲 De	evices			
INSPECTION	N TYPE	E					
□ New □ R	outine	☐ Cha	ange of Location 🗌 Ren	nodel 🗌 Other			
DESIGNATIO	ONS: C	MEA	NS COMPLIANT, NC M	IEANS NON-COMPLIANT			
FACILITY							
	C	NC					
54.1-3430			Permit is displayed in a	conspicuous place.			
110-20-630				rovided if operating from a privates during normal business hours.	te dwelling and the p	place of business is open	
SECURITY			for inspection at an time	s during normal business nours.			
	C	NC					
110-20-640	_	_	Areas in which prescription drugs are manufactured, stored, or kept for sale are held are restricted to only designated and necessary authorized personnel.				
110-20-640	_			shall provide reasonable security for			
110-20-640	_			shall not deliver any drug to a licer elivery nor to any person who may			
breaking, subj	ject to tl tested	he follo at time	wing conditions: of inspection	edical gases other than nitrous o	oxide, shall install a o	device for the detection of	
110-20-640			Device shall be a sound, device	microwave, photoelectric, ultrason	nic, or any other gene	rally accepted and suitable	
110-20-640	_	_		wired and both the installation and	l device shall be base	ed on accepted burglar	
110-20-640		_		ed in operating order and shall have	e an auxiliary source	of power	
110-20-640			Device shall fully protect breaking by any means w	t all areas where prescription drugs	s are stored and shall	be capable of detecting	
110-20-640	_		Access to the alarm syste	em shall be restricted to only design wer the drug storage areas are close		persons, and the system	
STORAGE							
	C	NC					
54.1-3461 (A)(3)			All prescription drugs as accordance with require	ormity with current good manufactive stored at appropriate temperature ments, if any, in the labeling of sumpendium, such as the United States	ures and under appro such drugs, or with re	equirements in the current	

RECORDS OF DRUGS IN SCHEDULES II, III, IV & V

	C	NC	
54.1-3404	_	_	Every inventory and other records required to be kept under this part shall be kept by the registrant and be available, for at least 2 years from the date of such inventory or records, for inspection and copying by authorized employees of the Administration, except that financial and shipping records (such as invoices and packing slips but not executed order forms subject to Sec. 1305.13 of this chapter) may be kept at a central location, rather than at the registered location, if the registrant has notified the Administration of his intention to keep central records. CFR 1304.04(a)
54.1-3404			The biennial inventory shall be taken on any date which is within two years of the previous biennial inventory.
54.1-3404	_	_	Date of inventory:
			☐ Opening or ☐ Closing of Business
54.1-3404	_	_	Inventories and records of Schedule II are maintained separately from all other records CFR 1304.04(f)(1)
54.1-3404	_	_	Inventories and records of controlled substances listed in Schedules III, IV, and V shall be maintained either separately from all other records of the registrant or in such form that the information required is readily retrievable from the ordinary business records of the registrant. CFR 1304.04(f)(2)
54.1-3404		_	The record of all drugs sold, administered, dispensed, or otherwise disposed of, shall show the date of selling, administering, or dispensing,
			 the name and address of the person to whom or for whose use, or the owner and species of animal for which the drugs were sold, administered or dispensed, kind and quantity of drugs.
			Any person selling, administering, dispensing or otherwise disposing of such drugs shall make and sign such record at the time of each transaction
54.1-3404		_	The record of such drugs received shall in every case show the • date of receipt, • name and address of the person from whom received • kind and quantity of drugs received, the kind and quantity of drugs produced or removed from process of manufacture
			 date of such production or removal from process of manufacture

Comments

This facility has been inspected by an inspector of the Department of Health Professions. The results of the inspection have been noted. I acknowledge that the noted conditions have been deemed by the inspector as not being in compliance and have been explained to me and that I have received a copy of the inspection report.

Signature of Inspector	Date	Signature of Licensee	Date